

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
23 August 2001 (23.08.2001)

PCT

(10) International Publication Number
WO 01/60325 A1

(51) International Patent Classification⁷: **A61K 7/04**,
A61F 13/00, A01N 35/00

(21) International Application Number: PCT/US01/05302

(22) International Filing Date: 16 February 2001 (16.02.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/182,924 16 February 2000 (16.02.2000) US

(71) Applicant (for all designated States except US):
BENTLEY PHARMACEUTICALS, INC. [US/US];
65 Lafayette Road, 3rd Floor, North Hampton, NH
03862-2403 (US).

(72) Inventor; and

(75) Inventor/Applicant (for US only): **GYURIK, Robert, J.**
[US/US]; 12 Ashbrook Road, Exeter, NH 03833 (US).

(74) Agent: **YAO, Gene, J.**; Synnestvedt & Lechner LLP, 2600
Aramark Tower, 1101 Market Street, Philadelphia, PA
19107-2950 (US).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: PHARMACEUTICAL COMPOSITION

(57) **Abstract:** A composition comprising a pharmaceutical compound, a membrane-permeation enhancer, and a polymeric film-forming agent, including, for example, a hydrocarbon-based liquid nail lacquer comprising a solution of an anti-fungal agent in a pharmaceutically-effective amount, a membrane-compatible permeation enhancer in an amount effective to enhance penetration through a membrane or nail of said anti-fungal agent, a polymeric film-forming agent in an amount effective to form an adherent polymeric film on a membrane or nail, and a solvent for forming said solution; and a method for forming on a membrane or nail of the body an adherent film containing a pharmaceutical compound and permeation enhancer from a solution of the components in which there is used a mixture of solvents, including a solvent which is highly volatile and a solvent of lower volatility.



WO 01/60325 A1

PHARMACEUTICAL COMPOSITION

Field of the Invention

The present invention relates to a composition useful for drug delivery.

- 5 More particularly, the present invention relates to a pharmaceutical composition which includes a permeation enhancer, that is, a material which is capable of increasing the rate of passage of a pharmaceutical compound through a nail and/or membrane.

The present invention will be described initially with respect to its use in the
10 treatment of fungal infections of the nail of the finger or toe. It should be understood, however, that the present invention can be used in other applications as well, as described below.

Onychomycosis is a fungal infection of the nail which affects 7 to 8 % of the North American population, including 15 to 20 % of adults aged 40 to 60 years and
15 25 to 40 % of adults over 60. Onychomycosis causes thickening and discoloration of the nail and can cause also loss or deterioration of the nail, pain, impaired circulation, and difficulty in walking, among other adverse effects.

Treatment of onychomycosis has in the past included measures such as removal of the infected portion of the nail or removal of the entire nail. This form

of treatment, however, may lead to permanent damage to the nail and the nail, if it grows back, may grow back in a misshapen form. In addition, there is no guarantee that onychomycosis is eradicated completely by nail removal.

Instead of nail removal, it is preferable to treat onychomycosis through the use of a variety of anti-fungal agents. Traditionally, such anti-fungal agents are administered orally. Upon administration, the agent moves through the body where small amounts of the agent arrive at the target area and penetrate the nail via the nail matrix. Oral administration is disadvantageous, however, in that such administration requires a prolonged treatment of about 12 weeks for toenails and about 6 to about 8 weeks for fingernails. Such prolonged treatment increases the cost of the treatment and reduces patient compliance. In addition, oral treatment creates a risk of intoxication, gastrointestinal irritation, nausea, adverse drug-to-drug interactions, drug-induced rash, and other adverse side effects. Moreover, variable rates of absorption and metabolism are often encountered in oral treatment.

Another method of treating onychomycosis involves the topical administration of a composition containing an anti-fungal agent. It is this method of treatment to which the present invention relates.

Reported Developments

It is known to treat the fungal infection of the nail by applying to the nail a pharmaceutical composition which contains an anti-fungal agent, for example, clotrimazole. The pharmaceutical composition is typically in the form of a nail lacquer, for example, as described in U.S. Patent No. 5,264,206. Such lacquers, however, have had limited success in the treatment of onychomycosis because the anti-fungal agents are not particularly effective in penetrating the nail and the skin surrounding and underneath the nail (the "nail bed").

The addition to the pharmaceutical composition of a permeation enhancer to improve the delivery of the anti-fungal drug to the target area has been proposed. For example, U.S. Patent No. 5,346,692 discloses a nail lacquer composition which comprises an anti-fungal agent and urea as a permeation enhancer. Urea functions,
5 however, by breaking down and disrupting the nail or skin in order to permit the anti-fungal agent to penetrate the deeper layers of the nail and nail bed. Accordingly, the use of urea results in damage to the nail and membrane.

Another composition for treating onychomycosis by applying the composition to the nail is disclosed in U.S. Patent No. 5,814,305. This composition
10 comprises: (A) an anti-fungal agent which is at least partially soluble in water; (B) a hydrophilic permeation enhancer; (C) water; and (D) alcohol. Anti-fungal agents of the allylamine family, such as terbinafine hydrochloride and naftifine hydrochloride, are preferred for use in the compositions of the '305 patent. The patent discloses the use of the following hydrophilic permeation enhancers: glycols; glycol diethers;
15 caprolactam; dimethylisosorbide; isopropylidene glycerol; dimethylimidazolidinone; N-methyl-pyrrolidone-2; pyrrolidone-2; ethyl lactate; and polyoxyethylenated glycerides. A disadvantage associated with the use of the composition described in the '305 patent is that a hydrophilic delivery system is used and most anti-fungal agents, in particular the azoles, are lipophilic. Lipophilic
20 enhancers have been shown to be more efficient than hydrophilic enhancers in increasing the absorption of pharmaceutical compounds into the strata of the membrane and the nail. Moreover, lipophilic antifungal agents are preferred since they have a tendency to penetrate more readily the membrane.

The present invention provides an improved composition that can be applied
25 to the nail for the purpose of treating a fungal infection that afflicts the nail and or nail bed and that can be used in other applications also.

Summary of the Invention

In accordance with this invention, there is provided a composition comprising: (A) a pharmaceutical compound; (B) a membrane-compatible permeation enhancer; and (C) a polymeric film-forming agent.

5 In addition, the present invention provides also a composition which is particularly suited to treating a nail disorder and which comprises a hydrocarbon-based liquid nail lacquer comprising a solution of: (A) an anti-fungal agent in a pharmaceutically-effective amount; (B) a membrane-compatible permeation enhancer in an amount effective to enhance penetration through a membrane or nail
10 of said anti-fungal agent; (C) a polymeric film-forming agent in an amount effective to form an adherent polymeric film on a membrane or nail; and (D) solvent for forming said solution.

Another aspect of the present invention is the provision of a method for administering to the body through the nail or membrane of the body comprising: (A)
15 applying to the nail or membrane a liquid composition comprising a pharmaceutical compound, a membrane-compatible permeation enhancer, and a polymeric film-forming agent; (B) forming from said liquid composition a solid film which adheres to the nail or membrane; and (C) maintaining said film on the nail or membrane for a period of time sufficient for delivery of the pharmaceutical compound to the body
20 through the nail or membrane.

Still another aspect of the present invention is the provision of a method for forming a solid adherent film comprising a pharmaceutical compound and an enhancer therefore comprising: (a) providing a solution in which the pharmaceutical compound, the enhancer, and a polymeric film-forming agent are dissolved, the
25 solution containing a co-solvent which includes: (i) a highly volatile solvent for the enhancer in which the enhancer is highly soluble; and (ii) a less volatile solvent in which the enhancer is less soluble; and (b) applying said soluble solution to the nail

or membrane; (c) forming from said solution a solid film which includes the pharmaceutical compound and the enhancer by permitting the highly volatile solvent to evaporate at a faster rate than the solvent having lower volatility; and (d) drying the resulting film.

- 5 It will be appreciated from the discussion which follows that the present invention provides an important advantage in that it can be used effectively to treat body conditions such as onychomycosis or other fungal infections, bacterial infections, and inflammation.

Detailed Description of the Invention

- 10 The composition of the present invention comprises a pharmaceutically effective amount of a pharmaceutical compound. Essentially any pharmaceutical compound which is capable of being delivered transdermally may be used in the practice of the present invention. The compound may be therapeutic or prophylactic. Examples of pharmaceutical compounds that may be employed in the
15 practice of the present invention include topically-effective agents, such as: anti-inflammatory corticosteroids; antibacterial agents; antiviral agents; and topical anti-cancer agents, such as 5-fluorouracil or other fluorinated purine, cytosine and pyrimidine analogs.

- In embodiments of the present invention in which anti-fungal agents are
20 used, the composition comprises a pharmaceutically effective amount of an anti-fungal agent which is capable of treating an infection residing in the nail and/or the nail bed. Essentially, any suitable anti-fungal agent may be employed. The anti-fungal agent may be present also in combination with an anti-inflammatory agent and/or another anti-microbial agent such as, for example, an anti-bacterial agent or
25 an anti-viral agent.

It is known that anti-fungal agents can function in various ways. For example, one class of anti-fungal agents works by impairing the functions of membrane-bound enzyme systems in the fungal cell membrane. Examples of such anti-fungal agents are azoles, for example, tioconazole, econazole, miconazole, 5 terconazole; clotrimazole, bifonazole, butaconazole, chlordantoin, chlormidazole, cloconazole, enilconazole, fenticonazole, isoconazole, ketoconazole, omoconazole, oxiconazole nitrate, and sulconazole.

It is known also that another class of anti-fungal agents functions by inhibition of oxidosqualene cyclase, thus deterring or preventing the formation of 10 the requisite ergosterol. Examples of such agents are allylamines, for example, terbinafine and naftifine.

Still another class of anti-fungal agents functions by compromising the integrity of the fungal cell membrane; an example of such an anti-fungal agent is ciclopirox.

15 Anti-fungal agents that work by varied modes of action as fungicidal or fungistatic agents may be employed also, for example, amorolfine, griseofluvin, nystatin, amphotericin B.

Clotrimazole and fluconazole are preferred anti-fungal agents for use in the practice of the present invention.

20 The pharmaceutical compound is present in the composition in a pharmaceutically effective concentration, which may be determined by those of ordinary skill in the art. Preferably, the concentration does not exceed the maximum amount that remains soluble in the composition, a parameter which can be determined readily also. For guideline purposes, it is believed most applications 25 will involve the use of the pharmaceutical compound in an amount of about 0.1% to about 15% by weight of the composition. A preferred amount of the compound is

about 1% to about 10% by weight of the composition, more preferably about 1% to about 8% by weight of the composition, and most preferably about 2% to about 5% by weight of the composition.

The composition comprises also a membrane-compatible permeation
5 enhancer which is capable of increasing the rate of passage of the pharmaceutical compound through a nail and/or membrane, that is, a layer of body tissue, for example, skin. The term "membrane-compatible permeation enhancer" means a compound which increases the rate of delivery of the pharmaceutical compound through the nail/membrane without damage.

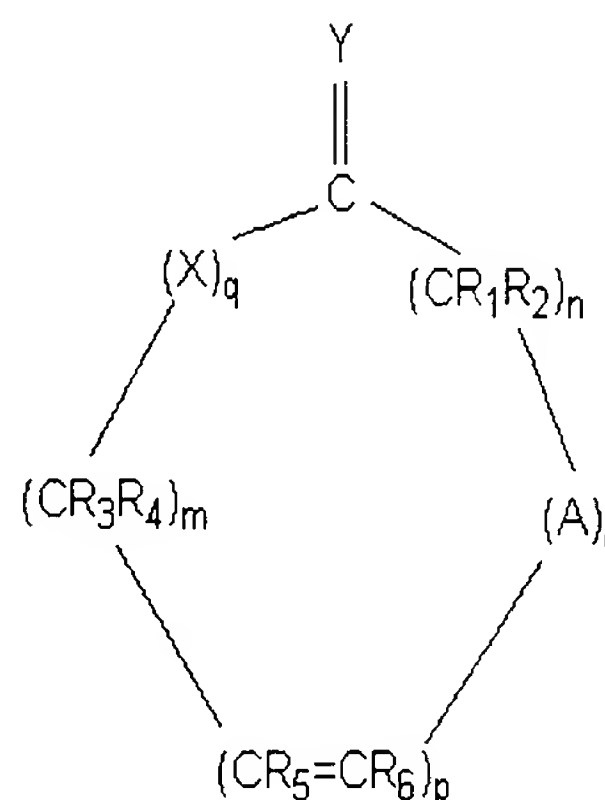
10 It is known that other enhancers function by a mechanism which involves hydrolysis, keratolysis, denaturation, or other mechanism which tends to damage the nail or the membrane. Examples of such enhancers include urea, sulfhydryl group-containing amino acids, alkyl sulfoxides, and related compounds which function by breaking down, extracting, or disrupting the nail or membrane in order to permit the
15 pharmaceutical compound to penetrate to deeper layers of the membrane.

The membrane-compatible enhancer of the present invention can be applied safely to the membrane or to a nail without causing damage to the membrane or nail.

Essentially any membrane-compatible permeation enhancer or mixture of
20 enhancers may be used in the practice of the present invention. Preferred membrane-compatible enhancers are lipophilic enhancers which render cellular membranes more permeable by intercalating within the membrane. Examples of such lipophilic enhancers include alkylesters, for example, isopropyl myristate and myristyl myristate.

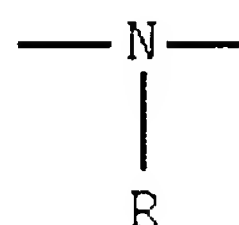
25 Examples of a particularly preferred class of membrane-compatible permeation enhancers are compounds which are described in U.S. Patent No.

5,023,252 to Hsieh (assigned to the same assignee as the present invention) and which have the structure of Formula I.

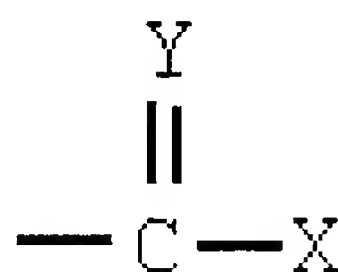


Formula I

5 wherein X and Y are oxygen, sulfur or an imino group of the structure



or =N-R, with the proviso that when Y is an imino group, X is an imino group, and when Y is sulfur, X is sulfur or an imino group, A is a group having the structure



10 wherein X and Y are as defined above, m and n are integers having a value from 1 to 20 and the sum of m+n is not greater than 25, p is an integer having a value of 0

or 1, q is an integer having a value of 0 or 1, r is an integer having a value of 0 or 1, and each of R, R₁, R₂, R₃, R₄, R₅, and R₆ is independently hydrogen or an alkyl group having from 1 to 6 carbon atoms which may be straight chained or branched, provided that only one of R₁ to R₆ can be alkyl group, with the proviso that when p, 5 q and r are 0 and Y is oxygen, then m+n is at least 11, and with the further proviso that when X is an imino group, q is equal to 1, Y is oxygen, and p and r are 0, then m+n is at least 11. For convenience, an enhancer of Formula I above is referred to herein as a "macrocyclic enhancer".

Macrocyclic enhancers have several desirable properties in addition to their 10 "enhancing" properties. For example, they have the ability to form a stable homogenous solution with the other components of the composition of the present invention and to function as a plasticizer and fluidizer in a film which is formed from the composition. This improves the performance properties of the film by rendering it pliable. This is surprising, because the preferred enhancer compounds, 15 for example, oxacyclohexadecan-2-one, are solids at room temperature. However, a liquid composition of the present invention in which the enhancer is dissolved is capable of forming a solid polymeric film which is clear and elastic. In addition, the film is capable of being resolubilized by successive applications of the film, providing a continuous, replenished delivery system without discontinuity or 20 disruption of the pharmaceutical compound.

Those skilled in the art will recognize which permeation enhancers are used preferably with the particular pharmaceutical compounds that are included in the composition. Particularly preferred membrane-compatible permeation enhancer for use in the practice of the present invention are oxacyclohexadecan-2-one, muscove, 25 civetone, and normuscione.

The membrane-compatible permeation enhancer is present in the composition in a concentration effective to enhance penetration through the nail and/or membrane of the pharmaceutical compound to be delivered. The "effective

amount” can be determined readily by the skilled artisan. For guideline purposes, it is believed most applications will involve the use of the membrane-compatible permeation enhancer in an amount of about 1% to about 25% by weight of the composition, preferably about 5% to about 20% by weight of the composition.

5 The composition comprises also a polymeric film-forming agent or mixture of film-forming agents. Essentially, there can be used any polymer that is capable of forming a film from which the pharmaceutical compound can be delivered to the nail or membrane, for example, the nail bed. For example, occlusive and semi-occlusive polymers which are known for use in transdermal drug delivery can be
10 used. Because the primary objective is drug delivery, even those polymers otherwise rejected for cosmetic end uses for less than ideal cosmetic properties can be used in the practice of the present invention.

 Examples of suitable polymeric film-forming agents include polymers of acrylic acid, acrylic acid esters and copolymers thereof; polymers of methacrylic
15 acid, methacrylic acid esters and copolymers thereof; polymers of vinyl acetate and copolymers thereof with acrylic acid and acrylic acid esters; copolymers of methyl vinyl ethers with maleic acid, maleic acid alkyl esters, and combinations thereof; copolymers of vinyl pyrrolidone with styrene; poly(vinyl butyrates); polymeric cellulose derivatives such as cellulose acetate phthalate, cellulose acetate butyrate,
20 cellulose acetate propionate, cellulose nitrate, cellulose sulfate, ethyl cellulose, and cellulose acetate; terpolymers of vinyl acetate with butyl maleate and isobornyl acetate; terpolymers of vinyl caprolactam with vinyl pyrrolidone and dimethylamino ethyl methacrylate. The film-forming agent can be used in solid, for example, powdery form in formulating a composition of the present invention. In addition,
25 the composition may be formulated also by use of a latex.

 Examples of preferred polymeric film-forming agents include: quaternary ammonium-containing acrylic acid ester; methacrylic acid ester copolymers known as ammonio methacrylate copolymers such as, for example, ethyl acrylate-[2-

(methacryloyloxy) ethyl] trimethylammonium chloride-methyl methacrylate copolymer; and substituted copolymers of alkylated poly(vinyl pyrrolidones). These polymeric film-forming agents are preferred because they demonstrate superior adhesive, water- resistance and hardness properties. Particularly preferred
5 are polymeric film-forming agents that have been registered with regulatory agencies for pharmaceutical use, including, but not restricted to the European or United States Pharmacopeia, and the Japanese Pharmaceutical Excipients Compendia.

The polymeric film-forming agent is present in an amount effective to form
10 on the membrane or the nail an adherent polymeric film. Amounts can be determined readily for any particular application. For guideline purposes, it is believed most applications will involve the use of the polymeric film-forming agent in an amount of about 0.1% to about 35%, preferably about 5% to about 35% by weight of the composition and more typically, and most preferably, in an amount of
15 about 10% to about 25% by weight of the composition.

The composition comprises also a solvent for those ingredients to be liquified. Essentially any solvent or solvent combination which is a suitable vehicle for the composition of the present invention can be employed. Examples of such solvents are alcohols, esters, ethers, aromatic hydrocarbons, aldehydes, ketones,
20 mono-, di- and tri-glycerides, and the like.

It is believed that the solvents that will be used most widely in the practice of the present invention are ethanol, ethyl acetate, butyl acetate, isopropanol, acetone, methyl ethyl ketone, triacetin, tripropionin, diethylene glycol monoethyl ether, and isopropyl acetate, and a mixture of two or more of the aforementioned. Ethanol,
25 ethyl acetate, butyl acetate, isopropanol, methyl ethyl ketone and acetone are preferred solvents as they evaporate and dry readily when applied.

The solvent is present in the composition in an amount sufficient to solubilize the ingredients that are to be liquified, for example, a solid polymeric film-forming agent, and other ingredients, without resulting in unsatisfactory drying times or film properties. For guideline purposes, it is believed most applications
5 will involve the use of the solvent in an amount of about 30% to about 80% by weight of the composition and more typically, and preferably, in an amount of about 40% to about 70% by weight of the composition.

In forming preferred compositions within the scope of the present invention, there are several factors that should be taken in to account in selecting the particular
10 pharmaceutical compound, enhancer, film-forming agent, and solvent that comprises the composition. The preferred composition is one which is capable of forming a solid film in a manner such that the concentrations of the pharmaceutical compound and enhancer are relatively high in the portion of the solid film which is contiguous to the surface on which the solid film is formed, for example, a nail
15 surface.

By way of background, it is noted that the term "fugacity" is used to refer to the measure of the escaping tendency of a solute in a solution and that the fugacity of a solute follows Henry's Law for ideal states. Various methods can be used in order to increase the fugacity of the pharmaceutical compound and enhancer, that is,
20 to increase the concentrations thereof at the desired site.

For example, with respect to a pharmaceutical compound which is basic in nature, the polymeric film-forming agent used in combination therewith is preferably also basic in nature. Such basic compounds tend to repel each because of the positive charges associated therewith, thus increasing the fugacity of the
25 pharmaceutical compound. Examples of polymeric film-forming agents that have a basic moiety or functionality are acrylate copolymers, for example, those that have intermittently distributed dimethylamino functionalities. Another method that can be used to increase the fugacity of the pharmaceutical compound and the film-

forming agent is to add to the solution thereof a basic compound, for example, TRIS amino or triethanolamine.

With respect to the use of an acidic-type pharmaceutical compound, for example, an acidic anti-fungal agent such as ciclopirox, there can be used a
5 polymeric film-forming agent that is acidic in nature. Examples of film-forming agents that have acid functionality are polymers or co-polymers of acrylic acid or methacrylic acid, for example, co-polymers of such acids with esters thereof.

The following is a description of a method to increase the fugacity of the permeation enhancer. The method involves formulating a composition comprising a
10 permeation enhancer dissolved in a volatile solvent, for example, a solvent which has a vapor pressure sufficient to allow it to evaporate within 5 minutes after application, and a less volatile solvent in which the permeation enhancer has limited solubility, for example, a solubility of up to 5 weight percent. Such a composition comprises co-solvents for the permeation enhancer. In the formation
15 of the film from the composition, the volatile solvent evaporates preferentially relative to the less-volatile solvent which remains (but temporarily) in the composition. As this occurs, the fugacity of the permeation enhancer increases. A cosolvent that increases the fugacity of a permeation enhancer upon the evaporation of a more volatile solvent in which the permeation enhancer is highly soluble
20 includes, for example, propylene glycol.

An exemplary method for increasing the fugacity of both the pharmaceutical compound, for example, an anti-fungal agent, which has limited solubility in water, and a permeation enhancer which has relatively high hydrophobicity (more limited water solubility than the pharmaceutical compound) is to include water in the
25 composition, for example, in an amount up to about 20% by weight of the composition, preferably about 1% to about 10%, and most preferably about 3% to about 7% by weight of the composition.

The foregoing description provides exemplary methods which may be employed to maximize the release of both the pharmaceutical compound and the enhancer or to provide more subtle increases and decreases in the fugacity of each component together or independently. Accordingly, the present invention provides
5 a means by which the release rate of either or both of the aforementioned components may be designed for more prolonged, controlled, or direct release of the medication.

One or more plasticizers can be included in the composition to impart desired properties to the film formed from the composition. In selecting a plasticizer and
10 amount to use, there should be taken into account whether the permeation enhancer that is present in the composition has plasticizing properties. Essentially any plasticizer can be employed. Examples of plasticizers are propylene glycol, diethylene glycol monoethyl ether, propylene glycol monopropyl ether, polyethylene and poly(propylene glycol), triacetin, tripropionin, castor oil, camphor,
15 phthalates, particularly dibutyl phthalate and diethyl phthalate, benzyl alcohol, phenethyl alcohol, and N-methyl-2-pyrrolidone, and a mixture of two or more of the aforementioned. As is known in the art, the plasticizer should be matched with the polymeric film-forming agent that is used in the composition.

It is believed that the plasticizers that will be used most widely in the practice
20 of the present invention will be propylene glycol, diethylene glycol monoethyl ether, polyethylene and poly(propylene glycol), triacetin, and tripropionin, and a mixture of two or more of the aforementioned. Propylene glycol, is among the preferred plasticizers.

The plasticizer is present in an amount sufficient to provide the desired
25 plasticizing properties to the polymeric film that is formed from the composition. For guideline purposes, it is believed most applications will involve the use of the plasticizer in an amount of 1% to about 25% by weight of the composition and more typically in an amount of about 1% to about 10% by weight of the composition.

The composition may include also other art-recognized components in art-recognized quantities. A coloring agent may be used, for example, a dye, color pigment, color lake, pearl gloss dye or pigment, for example, titanium dioxide, and the like. Other components include colloid stabilizers, UV stabilizers, antibacterial
5 or bacteriostatic substances such as quaternary ammonium antimicrobial agents, for example, cetyl pyridinium chloride, and benzalkonium chloride, anti-oxidants, for example, BHA, BHT, parabens, vitamin E and its derivatives, anti-microbial chelating agents, for example, EDTA and citric acid, and neutralizing agents, for example, TRIS amino, triethylamine, triethanolamine, 2-methyl-2-amino-1-
10 propanol, citric acid, and sorbic acid.

The composition of the present invention is applied to the membrane or the nail like any conventional composition that is capable of forming a solid film. The film can be formed from multiple coats of the composition. One or more successive coats may be applied typically after the underlying film is formed as the solvent
15 evaporates and the wet film dries. Periodic replacement of the film may be required to maintain the desired drug dosage regimen.

It is preferred that the components of the liquid composition be compatible with each other so that a clear film, free from clouding before and after application
20 results. The film should remain preferably clear and non-cloudy for each successive application. Clearness of the film is an indication that the desired molecules of anti-fungal agent and permeation enhancer are in a glassy state, and have not crystallized; otherwise the molecules may cease to migrate and not reach the intended target, the infected nail and membrane. Such a "cold flow" condition is
25 important to remediating the infection in an efficient manner.

As can be appreciated from the discussion above, the present invention provides an important advantage in that it can be used effectively to treat body conditions such as onychomycosis or other fungal infections, bacterial infections, and inflammation.

EXAMPLES

Examples below are illustrative of compositions of the present invention. The concentrations of the ingredients comprising the compositions are given in percent by weight relative to the total weight of composition.

- 5 Example Nos. 1 to 5 are examples of anti-fungal nail lacquers of the present invention. In the following examples, the polymeric film-forming agent, in powder or pelletized form, was dissolved in solvent while mechanically stirring at room temperature. After the film-forming agent was dissolved in the solvent, the pharmaceutical compound and the permeation enhancer, each in solid form, were
10 added with stirring. After the pharmaceutical compound and the permeation enhancer were dissolved, the plasticizer and water were added. The entire mixture was then stirred until homogenous.

Example No. 1

		<u>Wt. %</u>
15	clotrimazole, USP (Sifavitor)	4 %
	Eudragit RL 100 powder (Röhm) film-forming agent	15 %
	(ethyl acrylate, methyl methacrylate, trimethylammonioethyl methacrylate copolymer)	
	oxacyclohexadecan-2-one (Firmenich) - permeation enhancer	15 %
20	propylene glycol, USP - plasticizer	5 %
	ethanol, USP - solvent (200 proof)	58 %
	water	3 %

- Six (6) human patients with onychomycosis of the big toe, with at least 33% of the surface of the nail infected, were each treated once daily in the evening with
25 approximately 20-30 mg of a nail lacquer having the formulation of Example No. 1. The treatment was applied each day for seven days. The lacquer was removed on the seventh day of each week cycle by dissolving with 70% isopropyl alcohol.

Thereafter, the cycle was repeated for 120 days. After 60 days, substantial improvements were noted in all cases. Nails in all cases became clearer, harder, and the discoloration disappeared gradually. The cure proceeded from the cuticle to the distal nail. Complete cures, indicated by completely normal appearing nails, were achieved in all cases within 120 days of the beginning of the treatment. In two of the patients who were experiencing pain and tenderness in the nail, surprisingly, within one month after beginning treatment, the pain and tenderness subsided.

Example No. 2

10		<u>Wt. %</u>
	fluconazole (Quimica Sintética, SA)	2 %
	PVP/VA S-630 (ISP) powder film-forming agent (vinylpyrrolidone/vinyl acetate copolymer)	18%
	oxacyclohexadecan-2-one (Firmenich) - permeation enhancer	14 %
15	propylene glycol - plasticizer	4 %
	ethanol, USP - solvent (200 proof)	57 %
	tripropionin - plasticizer	1 %
	methyl-2-amino-1-propanol - neutralizing-agent	1%
	water	3 %

20 Example No. 3

		<u>Wt. %</u>
	clotrimazole (Sifavitor)	4 %
	Eudragit RL PO (Röhm) pellets - film-forming agent (ethyl acrylate, methyl methacrylate, tri- methylammonioethyl methacrylate copolymer)	15%
25	oxacyclohexadecan-2-one (Firmenich) - permeation enhancer	15%
	propylene glycol, USP - plasticizer	5 %
	ethanol, USP - solvent (200 proof)	58%
	water	3 %

Example No. 4

		<u>Wt. %</u>
	terconazole (Quimica Sintética, SA)	4%
	Gantrez® MS-955 (ISP) powder - film-forming agent	15 %
5	(2-butenedioic acid monobutyl ester, methoxyethylene copolymer)	
	cyclopentadecanone (Firmenich) - permeation enhancer	5%
	polyethylene glycol (MW 400) - plasticizer	5 %
	ethanol, USP - solvent (200 proof)	58 %
10	water	3 %

Example 5

		<u>Wt. %</u>
	fluconazole (Quimica Sintética, SA)	5%
15	Povidern™ SK3 (ISP) powder - film-forming agent (2-pyrrolidone, 1-ethenyl-, homopolymer)	15%
	Oxacyclohexadecan-2-one (Firmenich) - permeation enhancer	15%
	polyethylene glycol (USP) - plasticizer	10 %
	ethanol, USP (190 proof) - solvent	55 %
20	water	3 %

The next example is illustrative of a composition of the present invention in gel form.

Example No. 6

		<u>Wt. %</u>
	clotrimazole	1%
5	Carbopol 980 NF (BF Goodrich) powder	3%
	- film-forming agent and thickening agent	
	oxacyclohexadecan-2-one (Firmenich) - permeation enhancer	4%
	propylene glycol - plasticizer	5 %
	glycerin - plasticizer and humectant	3%
10	ethanol, USP - solvent (200 proof)	66 %
	triethanolamine, NF - neutralizing agent	Qs. to pH 5.5
	water, reagent grade	17 %

The gel of Example No. 6 was applied to excised guinea pig skin in a standard Franz cell apparatus (in vitro) or as a film on a 10 cm² area on live guinea pig skin (30 mg formulation applied) that had been prepared previously by shaving (in vivo). *In vitro* permeation time varied among 24, 48, and 72 hours and the total skin section was assayed. *In vivo* permeation time was seven (7) hours, after which the animals were euthanized, the skin washed, and skin separated into epidermis and dermis. The clotrimazole levels were measured in all cases by high performance liquid chromatography (HPLC). Lotrimin cream (1% clotrimazole) served as the control. The test results showed clearly a great magnitude of increase of permeation through the skin either *in vitro* or *in vivo* of the antifungal agent, clotrimazole, by one of the enhancing agents of the present invention, oxacyclohexadecan-2-one. It is believed that the present invention provides improved means for efficient and effective delivery of a pharmaceutical compound to the body by delivery thereof through a nail or membrane of the body.

CLAIMS

1. A composition comprising: (A) a pharmaceutical compound; (B) a membrane-compatible permeation enhancer; and (C) a polymeric film-forming agent.
- 5 2. A composition according to Claim 1 wherein said pharmaceutical compound is selected from the group consisting of: anti-fungal agents; estrogen; progesterone; nicotine; blood pressure-lowering agents; cholesterol-lowering agents; anti-depressant agents; nutritional supplements; and anti-dermatitis agents.
- 10 3. A composition according to Claim 1 wherein said pharmaceutical compound is an anti-fungal agent.
4. A composition according to Claim 3 wherein said anti-fungal agent is an azole.
- 15 5. A composition according to Claim 3 wherein said anti-fungal agent is clotrimazole.
6. A composition according to Claim 1 wherein said membrane-compatible permeation enhancer is lipophilic.
7. A composition according to Claim 1 wherein said membrane-compatible permeation enhancer is a macrocyclic enhancer.
- 20 8. A composition according to Claim 7 wherein said macrocyclic enhancer is oxacyclohexadecan-2-one.

9. A composition according to Claim 1 wherein said polymeric film-forming agent is selected from the group consisting of an: ammonio methacrylate copolymer; and a substituted copolymer of an alkylated poly(vinyl pyrrolidone).
10. A composition according to Claim 1 comprising also an anti-inflammatory compound.
11. A composition according to Claim 1 comprising also a solvent.
12. A composition according to Claim 1 wherein the solvent is selected from the group consisting of: ethanol; ethyl acetate; butyl acetate; isopropanol; acetone; methyl ethyl ketone; triacetin; tripropionin; diethylene glycol monoethyl ether; and isopropyl acetate; and a mixture of two or more of said solvents.
13. A composition according to Claim 1 comprising also a plasticizer.
14. A composition according to Claim 1 comprising from about 0.1 to about 15 wt. % of said pharmaceutical compound, from about 1 to about 25 wt. % said membrane-compatible permeation enhancer, and from about 5 to about 35 wt. % of said polymeric film-forming agent.
15. A hydrocarbon-based liquid nail lacquer comprising a solution of: (A) an anti-fungal agent in a pharmaceutically-effective amount; (B) a membrane-compatible permeation enhancer in an amount effective to enhance penetration through a membrane or nail of said anti-fungal agent; (C) a polymeric film-forming agent in an amount effective to form an adherent polymeric film on a membrane or nail; and (D) solvent for forming said solution.
16. A nail lacquer according to Claim 15 including about 0.1% to about 15% by weight of said anti-fungal agent; about 1% to about 25% by weight of said enhancer; about 0.1% to about 35% by weight of said film-forming agent; about

30% to about 80% by weight of said solvent; and a plasticizer in an amount of about 1% to about 25% by weight.

17. A method for administering a pharmaceutical compound to the body through the nail or membrane of the body comprising: (A) applying to the nail
5 or membrane a liquid composition comprising a pharmaceutical compound, a membrane-compatible permeation enhancer, and a polymeric film-forming agent; (B) forming from said liquid composition a solid film which adheres to the nail or membrane; and (C) maintaining said film on the nail or membrane for a period of time sufficient for delivery of the pharmaceutical compound to the body through the
10 nail or membrane.

18. A membrane according to Claim 17 wherein the pharmaceutical compound is an anti-fungal agent.

19. A method for forming on a membrane or a nail solid adherent film comprising a pharmaceutical compound and an enhancer therefore comprising: (A)
15 providing a solution in which the pharmaceutical compound, the enhancer, and a polymeric film-forming agent are dissolved, the solution containing a co-solvent which includes: (i) a highly volatile solvent for the enhancer in which the enhancer is highly soluble; and (ii) a less volatile solvent in which the enhancer is less soluble; and (B) applying said soluble solution to the nail or membrane; (C) forming
20 from said solution a solid film which includes the pharmaceutical compound and the enhancer by permitting the highly volatile solvent to evaporate at a faster rate than the solvent having lower volatility; and (D) drying the resulting film.

20. A method according to Claim 19 wherein said pharmaceutical compound is an anti-fungal agent and the solution is applied to a nail.

25 21. A composition according to Claim 1 wherein said pharmaceutical

compound, said enhancer, and said polymeric film-forming agent are dissolved in solution.

22. A composition according to Claim 21 wherein said solution comprises: (i) a highly volatile solvent for said enhancer in which said enhancer is highly soluble; and (ii) a less volatile solvent in which said enhancer is less soluble.

23. A composition according to Claim 1 wherein said pharmaceutical compound and said polymeric film-forming agent are basic in nature.

24. A composition according to Claim 23 wherein said composition comprises also a basic compound.

25. A composition according to Claim 24 wherein said basic compound is selected from the group consisting of TRIS amino and triethanolamine.

26. A composition according to Claim 1 wherein said pharmaceutical compound and said polymeric film-forming agent are acidic in nature.

27. A composition according to Claim 21 including also water.

28. A composition according to Claim 27 wherein water is present in an amount up to about 20 wt. % of the composition.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US01/05302

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61K 7/04; A61F 13/00; A01N 35/00

US CL : 424/61, 434; 514/675

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 424/61, 434; 514/675

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

NONE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 5,023,252 A (HSEIH) 11 JUNE 1991, see entire document.	1-28

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

21 MAY 2001

Date of mailing of the international search report

27 JUN 2001

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

SHARON HOWARD

Telephone No. (703) 308-1235

TERRY J. DEY

PARALEGAL SPECIALIST
TECHNOLOGY CENTER 1600